EXHIBIT 87

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Order Monitoring System (OMS): A Manufacturer's Perspective

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Mission of the Purdue OMS Program

To ensure compliance with DEA regulations requiring manufacturers and distributors to monitor and report suspicious orders of controlled substances, by implementing a detailed process for:

- Ongoing assessment of selected accounts, including Purdue's authorized distributors and their retail customers
- Support for authorized distributors in implementing their OMS programs and efforts to "know their customers"
- Reporting of suspicious ordering to DEA, other law enforcement, or state licensing boards, as appropriate

History of the Purdue OMS Program

- Followed DEA correspondence to all registrants detailing obligations of manufacturers and distributors of controlled substances to:
 - Conduct independent analysis and exercise due diligence to confirm legitimacy of orders and to scrutinize suspicious circumstances
 - Valid DEA registration not sufficient
 - Know your customers and your customers' customers
 - Inform DEA of suspicious orders when discovered
- Expanded program launched in 2008
- SOP finalized in March 2009

OMS Program Team Members

OMS COMMITTEE CHAIRPERSON

VP & Associate General Counsel, Law Department

MEMBERS

VP, Corporate Security
Executive Director, DEA Compliance
Executive Director, National Accounts
Director, OMS Program Coordinator
Director/Investigations, Corporate Security

CONTRIBUTORS

VP, Health Policy
Attorney, Prescriber Program analysis
Professional Rep, Sales Force
Director, Sales Systems

OMS Information Sources

- Fee For Service (FFS) Data
 - Order data for pharmacies + other dispensing outlets
 - Provided by authorized distributors under FFS Agreements
 - Loaded on monthly basis into OMS Database
 - Cover 97% of Purdue's product distribution
- IMS outlet/prescriber data & Sales Ops outlier analyses
- Sales Force reports of concern (ROC)
- Prescriber Program information
- Government agencies/law enforcement
 - DEA, local law enforcement, state licensing boards, legislative contacts
- Media reports

Prescriber versus Dispenser

- Prescriber program: Focus is on prescriber and Rx history /patterns
- OMS: Focus is on dispenser/pharmacist and ordering history/patterns
- Sharing of signal detection information between OMS and Prescriber programs
 - Enables us to consider prescriber and pharmacy issues within particular geographic area
 - Results in more robust information shared with internal (e.g., Risk Management) and external (e.g., authorized distributors) partners

OMS Process

- Identification of Potential Problematic Outlets ('09-'10)
 - FFS Data Outliers Outlets with orders outside normal range based on algorithm:
 - Total volume of Purdue product orders
 - Percentage of OxyContin / non-OxyContin orders to total orders of Purdue products
 - Percentage of orders of higher dosages of OxyContin
 - Number of distributors from which outlet purchases
 - Number of orders of same product per day
 - Significant increases/changes comparing current 1, 3, 6 and
 12 months to prior period

Based on algorithm, 500-600 outlets met criteria

OMS Process

(continued)

Identification of Problematic Outlets (continued)

- IMS Data Outliers
 - Outliers among retail outlets identified by Sales Ops' quarterly analysis of IMS Data
- Outlets identified by other signals
 - Typically identified by sales force or authorized distributors
 - Suspicious signals include:
 - Observed anomalies of pharmacy location, appearance/operation or clientele
 - Statements by pharmacy personnel indicating deficiencies in Rx verification or other abuse/diversion mitigation procedures
 - Authorized distributor comparative data on other opioid dispensing by pharmacy or Rx detail on pharmacy's prescribers
 - Media reports of law enforcement or licensing board action

- Outlier Pharmacies Selected for Review:
 - Top FFS Data Outliers (as ranked by Sales Ops)
 - Accounts identified by authorized distributors
- Input from National Accounts
 - Any prior knowledge of pharmacy, including factors that explain or heighten concern about outlier data
 - Assessment of need for further follow up
- > Input from Sales Force
 - Review of prior ROCs
 - Standard OMS follow with Rep / DM / RM
 - Specific additional assistance occasionally requested

OMS Process

(continued)

- Review of Related Internal Data & Information
 - Savings Card Pharmacy Redemption data
 - Analysis of identified prescribers (IMS data)
- > Public Records Search
 - Corporate security review of entity status and ownership, including related entities
 - DEA registration / state licensing status and disciplinary actions
 - Civil or criminal actions

OMS Process

(continued)

- DEA Compliance: Collaboration with Authorized Distributors
 - Initial meetings to share information about respective order monitoring programs and procedures
 - Ongoing information exchange and review of ordering data and other information pertaining to specific outlets
 - Communication and collaboration on follow up with respect to individual outlets, which may include:
 - Outlet surveillance and/or site visit and interview of owner,
 Pharmacist-In-Charge and/or pharmacy staff
 - Reduction or cut-off of supply to outlet
 - Reporting to licensing board, DEA, other law enforcement

Summary of OMS Meetings

- Order monitoring meetings held with authorized distributors plus ongoing contacts:
 - Between Sept 2008 and March 2012, Purdue met in person with 10 separate wholesalers to discuss OMS programs and procedures, and opportunities for better collaboration
 - Throughout that time, Purdue engaged in regular ongoing contact via conference calls and joint site visits to discuss particular accounts of concern and appropriate follow up

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OMS Report and Committee Decision

- Written report for OMS Committee review
 - Generated by Program Coordinator for each "outlier outlet"
 - Captures information obtained during OMS review process
- OMS Committee decision on each outlet reviewed
 - Pending: No decision pending completion of requested follow up
 - Complete-closed: No suspicious ordering concern
 - Complete-referred: Evidence of suspicious ordering and/or circumstances sufficient to refer to DEA, other law enforcement and/or state licensing board
 - Continue to monitor: Suspicious circumstances warrant close monitoring, but not yet sufficient to refer
- OMS Committee may recommend adjustments in shipments to distributor due to OMS concerns

OMS Process: Post Reformulation

- Updated Algorithm Based on Reformulation ('10 '11)
 - FFS Data Outliers Outlets with decline in orders post OxyContin reformulation that met the following:
 - Orders that met original algorithm
 - Significant declines/changes comparing current 3, 6 and 12 months of pre- versus post-reformulation data
 - Threshold 75% decline post reformulation
 - Percentage of OxyContin decline post reformulation vs contemporaneous increase in other opioids
 - Evaluate whether geographically located near prescribers of concern

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Adjust threshold (\$) to focus on significant accounts for review

Based on new algorithm, 100 to 200 outlets met criteria

Meetings with DEA

≻April 2009

- Overview of OMS program
- Described collaboration with authorized distributors

≻April 2011

- Overview of updated Purdue OMS program following reformulation
- DEA Registrant book shared
- Focus on prescriber data post-reformulation

➢October 2011

- Focus on retail dispensing post-reformulation
- At request of DEA, provided calculation of all outlets with at least 50% decline and \$350,000 in annual sales
- Total of 290 outlets identified (29 previously identified)

Summary of OMS Program Activity

(continued)

Outlets Reviewed and/or Referred ('08 – '11)

• Total: 365

Breakdown by state:

FL: 94
 CA: 55
 NY: 39
 O PA: 18
 O TN: 14
 O OH: 13

o MI: 38 o 27 States: 94 (3 to 4 each)

Breakdown by OMS Committee Action:

Complete-Referred: 290

Complete-Closed: 75

Continue to Monitor: 8

Summary OMS Program Activity

(continued)

- Outlets pending review/investigation
 - **Total**: 8
 - Breakdown by state:
 - \circ GA = 3
 - NY/NJ/CA/TN/IN = 5 (1 each)
- Outlets subject to OMS Team Surveillance or Site Visits
 - 13 pharmacy site visits including interviews with owners or pharmacists in charge
 - 6 of the visits conducted together with authorized distributors
 - Breakdown by location: 8 in Florida, 2 in California and Nevada, 1 in NY
 - 10 additional pharmacies subject to surveillance
 - o 5 in California, 2 each in Ohio and Florida, 1 in Nevada
 - 30 + site visits with wholesalers

OMS Program Challenges

- Data Gaps
 - No data connecting outlets with individual prescribers
 - No data from distributors with whom we have no FFS agreement
 - FFS data excludes outlet-level order detail for:
 - Secondary distributors
 - Dispensing outlets that opt out of data reporting
 - IMS data excludes prescribers/outlets who opt out of reporting
 - Dispensing healthcare providers
- Not in doctors office, or at pharmacy, when prescriptions being written and filled
- Pressure Created by Geographic Hotspots (e.g., Florida, California, Tennessee, Georgia and Alabama)

Recommendations: Lessons Learned

- Quantities matter: excessive orders must be evaluated
- Meaningful scrutiny of dispensing: registration not sufficient
- Site visit due diligence: expected as part of follow up
- Cannot rely on third party: must do own due diligence
- Trend analysis is a key: compare similar products, size and location of outlets
- Threshold exceptions: must be individually reviewed and decisions properly documented
- Referrals to DEA: consider for all OMS actions regarding outlets

Benefits of Collaboration: What can be gained?

- > Enhance collaborations efforts between wholesalers and manufacturers
- Greater information sharing: maximize resources (DEA, Wholesaler and Manufacturer)
- Achieve efficiencies with accounts identified for follow up
- Identify additional tools to address DEA's concerns (better data analysis, potential modeling)
- Mindful of anti-trust concerns

Thank You

Any Questions

